

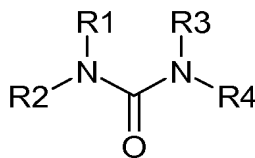
Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claims 1-16 (cancelled)

Claim 17 (currently amended): A separation matrix for affinity chromatography, comprising ligands coupled to a support, wherein the majority of the ligands are the compounds of formula (I)



(I)

wherein

R₁ is CH₃ or CH₂CH₃;

R₂ is a ~~*para* and/or *meta*~~ substituted-phenyl group that has been substituted with Cl in *meta* and *para* position having substituents selected from the group consisting of F, Cl, Br, I, OH and a group —O—R₅, wherein R₅ is either CH₂ or CH₂CH₃;

R₃ is H, CH₃ or CH₂CH₃; and

R₄ is a linear or cyclic aliphatic group,

or, wherein

R₁ and R₂ are as stated above while R₃ and R₄ are parts of a 4- to 6-membered cyclic entity,

and which compound has affinity for human IgG of κ -type;

further wherein said ligands are coupled to said support through the group R₄.

Claim 18 (previously presented): The separation matrix of claim 17, wherein the ligands have been coupled to the support via linkers.

Claim 19 (previously presented): The separation matrix of claim 17, wherein the support is a porous polymeric particle.

Claim 20 (cancelled)

Claim 21 (withdrawn): A system suitable for affinity chromatography, comprising the separation matrix of claim 17 packed in a column.

Claim 22 (previously presented): The separation matrix of claim 17, wherein the compounds of formula (I) is an affinity ligand with affinity for the constant region of a Fab fragment of human IgG of κ -type.

Claim 23 (previously presented): The separation matrix of claim 17, wherein R₁ is CH₃.

Claims 24-27 (cancelled)

Claim 28 (previously presented): The separation matrix of claim 17, wherein R₄ is an aliphatic group, which includes oxygen atoms in one or more positions.

Claim 29 (previously presented): The separation matrix of claim 17, wherein R₄ is an aliphatic group, which contains one or more carbonyl groups.

Claim 30 (cancelled)

Claim 31 (currently amended): The separation matrix of claim 17, wherein R₁ is CH₃; ~~R₂ is a phenyl group that has been substituted with Cl in *meta* and *para* position;~~ and R₃ and R₄ are parts of a cyclic 5-membered group.

Claim 32 (previously presented): The separation matrix of claim 31, wherein the cyclic 5-membered entity is substituted in a position directly adjacent to N with a C(O)-O-CH₃ group.

Claim 33 (previously presented): The separation matrix of claim 17, wherein said compounds of formula (I) are capable of binding to the constant region of a human

IgG of κ -type, or a functional derivative thereof, with a binding constant of at least 10^{-3} M.

Claim 34 (previously presented): The separation matrix of claim 17, wherein said compounds of formula (I) are capable of binding to the constant region of a human IgG of κ -type, or a functional derivative thereof, via a binding pocket-defined by the structure coordinates of the amino acids as shown in Fig 6.